

Linco-Spectin 100 222 mg/g - 444.7 mg/g Powder for use in drinking water

Authorised

- Spectinomycin sulfate
- Lincomycin hydrochloride

Product identification

Medicine name:

Linco-Spectin 100 222 mg/g - 444.7 mg/g Powder for use in drinking water

Active substance:

Spectinomycin sulfate

Lincomycin hydrochloride

Target species:

Pig

Chicken

Route of administration:

Oral use

Product details

Active substance and strength:

Spectinomycin sulfate

575.90 milligram(s) / 1.00 gram(s)

Lincomycin hydrochloride

241.90 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water

Withdrawal period by route of administration:

Oral use:

-

Pig

- Meat and offal. 0 day

Animals must not be slaughtered for human consumption during treatment.

-

Chicken

- Meat and offal. 5 day

- Egg. no withdrawal period

Not for use in birds producing or intended to produce eggs for human consumption

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FF52

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Available in:

Belgium

Package description:

Linco-Spectin 100 222 mg/g - 444.7 mg/g powdr. for drinking water 1.5 kg

Linco-Spectin 100 222 mg/g - 444.7 mg/g powdr. for drinking water 150 g

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Phibro Animal Health (Poland) Sp. z o.o.

Marketing authorisation date:

1/06/1972

Manufacturing sites for batch release:

Zoetis Belgium

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V061957

Date of authorisation status change:

7/07/2020

Reference member state:

Belgium

Procedure number:

BE/V/0029/001

Concerned member states:

Austria Cyprus Czechia Denmark Estonia France Germany Greece Hungary
Ireland Italy Latvia Lithuania Luxembourg Poland Portugal Slovakia Spain
United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Package Leaflet

English (PDF)

Published on: 24/03/2026

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Summary of Product Characteristics

English (PDF)

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Labelling

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English (PDF)

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